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Phase II trial of ZD1839 in recurrent or metastatic squamous cell car the head and neck.
J Clin Oncol. 2003 May 15;21(10):1980-7.
PMID: 12743152 [PubMed - indexed for MEDLINE]

2: Baselga J, Rischin D, Ranson M, Calvert H, Raymond E, Kieback DG, Kaye SB, Gianni L, Harris A, Bjork T, Averbuch SD, Feyereislova A, Swaisland H, Rojo F, Albanell J. Related

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J Clin Oncol. 2002 Nov 1;20(21):4292-302.
PMID: 12409327 [PubMed - indexed for MEDLINE]

3: Herbst RS, Maddox AM, Rothenberg ML, Small EJ, Rubin EH, Baselga J, Rojo F, Hong WK, Swaisland H, Averbuch SD, Ochs J, LoRusso PM. Related

Selective oral epidermal growth factor receptor tyrosine kinase inhib ZD1839 is generally well-tolerated and has activity in non-small-cell cancer and other solid tumors: results of a phase I trial.
J Clin Oncol. 2002 Sep 15;20(18):3815-25.
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OSI Pharmaceuticals, Genentech and Roche announce data from clin of Tarceva.
Expert Rev Anticancer Ther. 2001 Jun;1(1):4-5. No abstract available.
PMID: 12113131 [PubMed - indexed for MEDLINE]

5: Dittrich Ch, Greim G, Borner M, Weigang-Kohler K, Huisman H, Amelsberg A, Ehret A, Wanders J, Hanauske A, Fumoleau P. Related

Phase I and pharmacokinetic study of BIBX 1382 BS, an epidermal g factor receptor (EGFR) inhibitor, given in a continuous daily oral administration.
Eur J Cancer. 2002 May;38(8):1072-80.
PMID: 12008195 [PubMed - indexed for MEDLINE]

6: Ranson M, Hammond LA, Ferry D, Kris M, Tullo A, Murray PI, Miller V, Averbuch S, Ochs J, Morris C, Feyereislova A, Swaisland H, Rowinsky EK. Related

ZD1839, a selective oral epidermal growth factor receptor-tyrosine k inhibitor, is well tolerated and active in patients with solid, malignan results of a phase I trial.
J Clin Oncol. 2002 May 1;20(9):2240-50.
PMID: 11980995 [PubMed - indexed for MEDLINE]

7: Ellis MJ, Coop A, Singh B, Mauriac L, Llombert-Cussac A, Janicke F, Miller WR, Evans DB, Dugan M, Brady C, Quebe-Fehling E, Borgs M. Related

Letrozole is more effective neoadjuvant endocrine therapy than tamo ErbB-1- and/or ErbB-2-positive, estrogen receptor-positive primary l cancer: evidence from a phase III randomized trial.
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PMID: 11559718 [PubMed - indexed for MEDLINE]

8: Hidalgo M, Siu LL, Nemunaitis J, Rizzo J, Hammond LA, Takimoto C, Eckhardt SG, Tolcher A, Britten CD, Denis L, Ferrante K, Von Hoff DD, Silberman S, Rowinsky EK. Related

Phase I and pharmacologic study of OSI-774, an epidermal growth fa receptor tyrosine kinase inhibitor, in patients with advanced solid ma
J Clin Oncol. 2001 Jul 1;19(13):3267-79.
PMID: 11432895 [PubMed - indexed for MEDLINE]

9: Ghosh S, Liu XP, Zheng Y, Uckun FM. Related

Rational design of potent and selective EGFR tyrosine kinase inhibit anticancer agents.
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Curr Opin Investig Drugs. 2001 Feb;2(2):298-304. Review.
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11: Bridges AJ. Related

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Chem Rev. 2001 Aug;101(8):2541-72. Review. No abstract available.
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Tyrosine kinase inhibitors-ZD1839 (Iressa).
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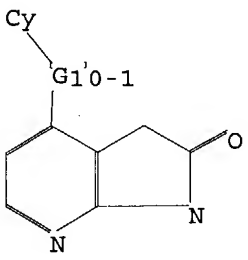

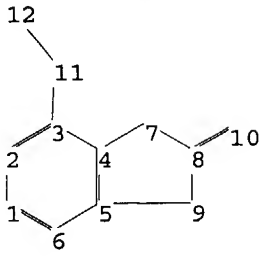

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ring nodes :
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ring bonds :
1-2 1-6 2-3 3-4 4-5 4-7 5-6 5-9 7-8 8-9
exact/norm bonds :
3-11 4-7 5-9 7-8 8-9 8-10 11-12

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exact bonds :
13-14
normalized bonds :
1-2  1-6  2-3  3-4  4-5  5-6


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     pyrrolo[2,3-b]pyridin-4-yl]phenyl]-6-(methylamino)- (9CI)  (CA INDEX NAME)
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SR   CA
LC   STN Files:   CA, CAPLUS, USPAT2, USPATFULL
DT.CA  CAplus document type:  Patent
RL.P   Roles from patents:  BIOL (Biological study); PREP (Preparation); USES
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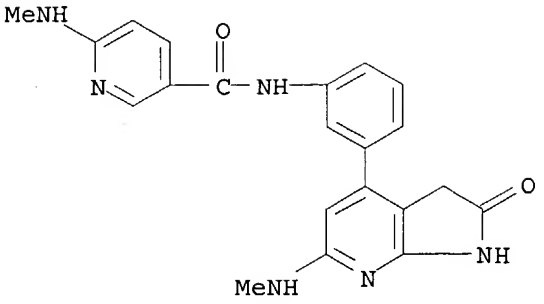

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)
1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

REFERENCE 1: 135:76886 Preparation of 4-substituted 7-azaindolin-2-ones and their use as protein kinase inhibitors. Liang, Congxin; Sun, Li; Wei, Chung Chen; Tang, Peng Cho; McMahon, Gerald; Hirth, Klaus Peter; Cui, Jingrong (Sugen, Inc., USA). PCT Int. Appl. WO 2001046196 A1 20010628, 97 pp. DESIGNATED STATES: W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG, TR. (English). CODEN: PIXXD2. APPLICATION: WO 2000-US34259 20001221. PRIORITY: US 1999-PV171288 19991221.

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	JP 2003518123	T2	20030603	JP 2001-547106	20001221
	US 2004053924	A1	20040318	US 2003-622787	20030721

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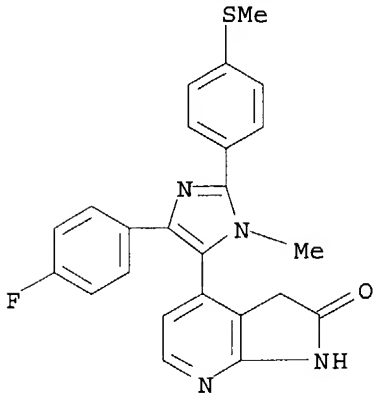
CN 2H-Pyrrolo[2,3-b]pyridin-2-one, 4-[4-(4-fluorophenyl)-1-methyl-2-[4-(methylthio)phenyl]-1H-imidazol-5-yl]-1,3-dihydro- (9CI) (CA INDEX NAME)
FS 3D CONCORD
MF C24 H19 F N4 O S
SR CA

LC STN Files: CA, CAPLUS, USPAT2, USPATFULL
DT.CA CAplus document type: Patent
RL.P Roles from patents: BIOL (Biological study); PREP (Preparation); USES (Uses)



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SR CA
LC STN Files: CA, CAPLUS, USPAT2, USPATFULL
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RL.P Roles from patents: BIOL (Biological study); PREP (Preparation); USES (Uses)

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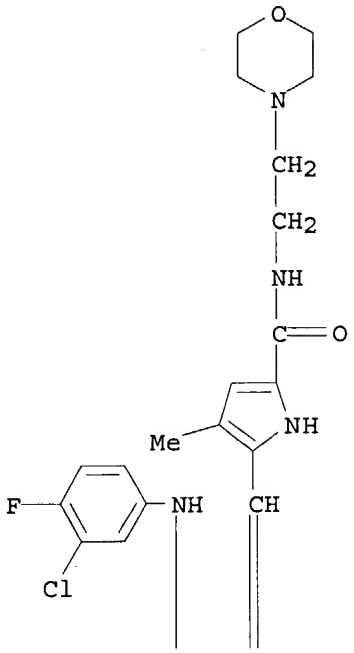

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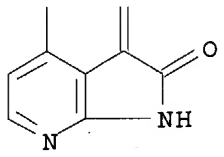

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